'M. M. Daicia 10/635808

=> dis his

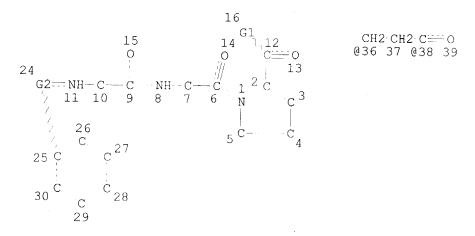
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DEL HIS Y

FILE 'REGISTRY' ENTERED AT 10:16:44 ON 28 SEP 2004
L1 STR
L2 50 S L1
L3 STR L1
L4 1 S L3
L5 STR L3
L6 308 S L5 FUL

=> d 16 que stat; fil medl, hcapl, embase, biosis; s 16 L5 STR

 Ak-NH
 O Ak
 Ak N-Ak
 C=0
 CH2-C=0

 @17 18
 @19 20
 21 @22 23
 @31 32
 @33 @34 35



CH2-CH2-CH2-C=-O @40 41 42 @43 44 CH CH C O 045 46 047 48 O-VCH2·C=O 049 50 051 52

VAR G1=OH/19/NH2/17/22 VAR G2=31/33-25 34-11/36-25 38-11/40-25 43-11/45-25 47-11/49-25 51-11 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 52

STEREO ATTRIBUTES: NONE
L6 308 SEA FILE=REGISTRY SSS FUL L5

100.0% PROCESSED 294255 ITERATIONS SEARCH TIME: 00.00.27

308 ANSWERS

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 166.34 690.09

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE ENTRY TOTAL SESSION -37.10

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TOTAL FOR ALL FILES L11 234 L6

=> s l11 and (postlesion? or post lesion? or neurodegenerat? or neurolog? degen? or alzheimer? or dement? or cognitive impairment or neural trauma)

L12 0 FILE MEDLINE
L13 3 FILE HCAPLUS
L14 0 FILE EMBASE
L15 0 FILE BIOSIS

TOTAL FOR ALL FILES

L16 3 L11 AND (POSTLESION? OR POST LESION? OR NEURODEGENERAT? OR NEURO LOG? DEGEN? OR ALZHEIMER? OR DEMENT? OR COGNITIVE IMPAIRMENT OR NEURAL TRAUMA)

=> d 1-3 cbib abs hitstr

L16 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2004 ACS on STN
2002:609522 Document No. 137:163818 Tripeptide derivatives for the treatment of post-lesional diseases of the nervous system.

Rapin, Jean; Witzmann, Hans Klaus; Grumel, Jean-Marie; Gonella, Jacques (Tell-Pharm AG, Switz.). Ger. Offen. DE 10105040 A1 20020814, 4 pp. (German). CODEN: GWXXBX. APPLICATION: DE 2001-10105040 20010205.

 R^{1}

0

 R^5 I

AB The invention discloses the use of cinnamoyl tripeptide derivs. for the

treatment of post-lesional neuronal diseases. The cinnamoyl tripeptide derivs. are I $[X=OH,\ C1-5\ alkoxy,\ NH2,\ NH(C1-5\ alkyl),\ N(C1-5\ alkyl)2;\ R=(preferably)\ cinnamoyl;\ R1=group\ derived from Phe, Tyr, Trp, Pro, Ala, Val, Leu or Ile; R2=group\ derived from Gly, Ala, Ile, Val, Ser, Thr, His, Arg, Lys, Pro, Glu, Gln, pGlu, Asp and Asn; R3, R4=H, OH, C1-5 alkyl, C1-5 alkoxy, provided that R3 and R4 are not both OH or C1-5 alkoxy; R5=H, OH, C1-5 alkyl, C1-5 alkoxy], or a pharmaceutical acceptable salt thereof.$

IT 123910-57-6

RN

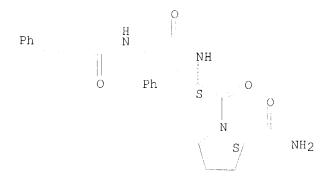
RL: PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (tripeptide derivs. for treatment of post-lesional nervous system diseases)

123910-57-6 HCAPLUS

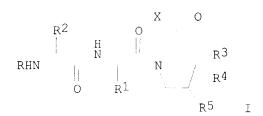
CN L-Prolinamide, N-(1-oxo-3-phenyl-2-propenyl)glycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry unknown.



L16 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2004 ACS on STN
2002:591566 Document No. 137:135103 Tripeptide derivatives for treatment of
neurodegenerative diseases. Rapin, Jean; Witzmann, Hans Klaus;
Grumel, Jean-Marie; Gonella, Jacques (Tell-Pharm A.-G., Switz.). Ger.
Offen. DE 10105039 A1 20020808, 10 pp. (German). CODEN: GWXXBX.
APPLICATION: DE 2001-10105039 20010205.



The invention discloses the use of tripeptide derivs. for treatment of neurodegenerative diseases. The tripeptide derivs. are I [X = OH, C1-5 alkoxy, NH2, NH(C1-5 alkyl), N(C1-5 alkyl)2; R = (preferably) cinnamoyl; R1 = group derived from Phe, Tyr, Trp, Pro, Ala, Val, Leu or Ile; R2 = group derived from Gly, Ala, Ile, Val, Ser, Thr, His, Arg, Lys,

Pro, Glu, Gln, pGlu, Asp or Asn; R3, R4 = H, OH, C1-5 alkyl, C1-5 alkoxy, provided that R3 and R4 are not both OH or C1-5 alkoxy; R5 = H, OH, C1-5 alkyl, C1-5 alkoxy], or a pharmaceutically compatible salt. Cinnamoyl-Gly-L-Phe-L-Pro-NH2 was tested in an Alzheimer's disease model.

ΙT 123910-57-6

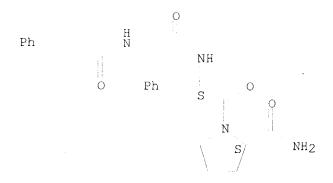
> RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (tripeptide derivs. for treatment of neurodegenerative

diseases)

RN 123910-57-6 HCAPLUS

L-Prolinamide, N-(1-oxo-3-phenyl-2-propenyl)glycyl-L-phenylalanyl- (9CI) CN(CA INDEX NAME)

Absolute stereochemistry. Double bond geometry unknown.



L16 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2004 ACS on STN 1989:633680 Document No. 111:233680 Preparation of tripeptides containing L-proline derivatives as nootropics and pharmaceutical compositions containing them. Fiez-Vandal, Pierre Yves (Inorgan S. A., Switz.). Eur. Pat. Appl. EP 316218 A1 19890517, 18 pp. DESIGNATED STATES: R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE. (French). CODEN: EPXXDW. APPLICATION: EP 1988-402761 19881103. PRIORITY: FR 1987-15228 19871103. GI

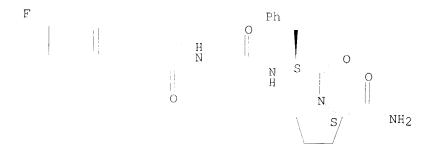
AΒ The title compds. [I; R1 = Q; X = CO, YCO, OYCO; Y = alkylene, alkenylene; Z = H, ≥1 CF3, alkyl, alkylenedioxy; R2 = NH2, OH, or a functional derivative thereof; A1, A2 = amino acid residue; B1, B2 = H, Me] and their pharmaceutically acceptable salts, useful as nootropics for treatment of senile dementia, Alzheimer's disease, Parkinson's disease, schizophrenia, and depression, are prepared via reaction of activated R1-A1-OH with proline derivs. II (R3 = H-A2), obtained by reaction of II (R3 = H) with activated H-A2-OH. N-Cinnamoylglycine (preparation given) was condensed with II.CF3CO2H (R2 = NH2, $B\bar{1}$ = B2 = H, R3 = H-Phe) (preparation given) in DMF containing dicyclohexylcarbodiimide and N-methylmorpholine to give I (R1 = cinnamoyl, R2 = NH2, B1 = B2 = H, A1 =

Gly, A2 = Phe) (III). III, administered i.p. or p.o. at 1 mg/kg, was effective in antagonizing scopolamine-induced amnesia in mice.

123910-50-9P 123910-52-1P 123910-53-2P 123910-54-3P 123910-55-4P 123910-56-5P 123910-57-6P 123910-58-7P RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of, as nootropic)

RN 123910-50-9 HCAPLUS
CN L-Prolinamide, N-[3-(4-fluorophenyl)-1-oxo-2-propenyl]glycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

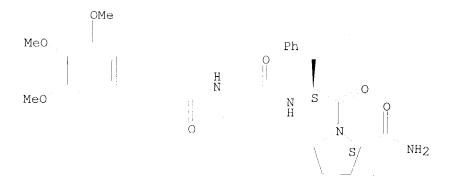
Absolute stereochemistry.
Double bond geometry unknown.



RN 123910-52-1 HCAPLUS
CN L-Prolinamide, N-[1-oxo-3-(3,4,5-trimethoxyphenyl)-2-propenyl]glycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

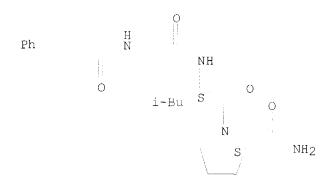
Absolute stereochemistry.

Double bond geometry unknown.



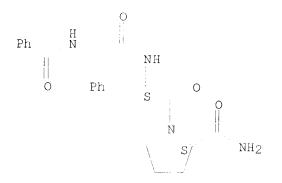
RN 123910-53-2 HCAPLUS
CN L-Prolinamide, N-(1-oxo-3-phenyl-2-propenyl)glycyl-L-leucyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



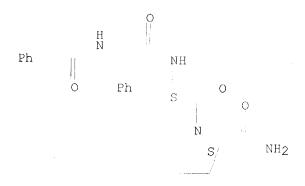
RN 123910-54-3 HCAPLUS CN L-Prolinamide, N-benzoylglycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



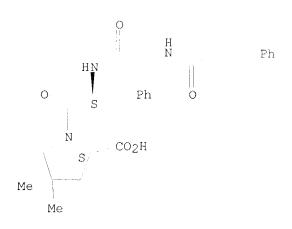
RN 123910-55-4 HCAPLUS
CN L-Prolinamide, N-(phenylacetyl)glycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



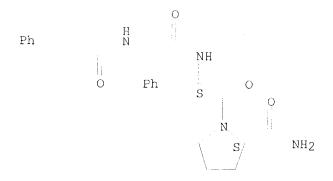
RN 123910-56-5 HCAPLUS
CN L-Proline, 4,4-dimethyl-1-[N-[N-(1-oxo-3-phenyl-2-propenyl)glycyl]-L-phenylalanyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry unknown.



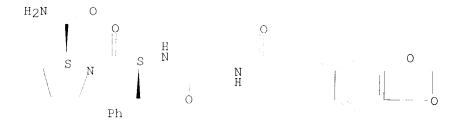
RN 123910-57-6 HCAPLUS
CN L-Prolinamide, N-(1-oxo-3-phenyl-2-propenyl)glycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.
Double bond geometry unknown.



RN 123910-58-7 HCAPLUS
CN L-Prolinamide, N-[3-(1,3-benzodioxol-5-yl)-1-oxo-2-propenyl]glycyl-L-phenylalanyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry unknown.



=> fil hcapl;e postlesional/ct COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 38.79 728.88

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE
ENTRY
SESSION
-2.10
-39.20

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FILE COVERS 1907 - 28 Sep 2004 VOL 141 ISS 14 FILE LAST UPDATED: 27 Sep 2004 (20040927/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

E#	FREQUENCY	AT	TERM
E1	0	1	POSTIMPLANTATION/CT
E2	0	2	POSTIMPLANTATION EMBRYO/CT
ЕЗ	0	>	POSTLESIONAL/CT
E4	0	1	POSTMENOPAUSAL/CT
E5	0	2	POSTMENOPAUSAL BONE LOSS/CT
E6	0	2	POSTMENOPAUSAL HORMONE REPLACEMENT THERAPY/CT
E7	0	2	POSTMENOPAUSAL OSTEOPOROSIS/CT
E8	0	2	POSTMENOPAUSE/CT
E9	0	1	POSTMORTEM/CT
E10	0	2	POSTMORTEM HUMAN TISSUE/CT
E11	0	1	POSTNATAL/CT
E12	0	2	POSTNATAL DEPRESSION/CT

=> e neurodegenerative/ct

Ε#	FREQUENCY	AT	TERM
E1	0	1	NEUROD3/CT
E2	0	2	NEURODEGENERATION/CT
ЕЗ	0	1	> NEURODEGENERATIVE/CT
E4	0	2	NEURODEGENERATIVE DISEASE/CT
E5	0	2	NEURODEGENERATIVE DISEASES/CT
E6	0	2	NEURODEGENERATIVE DISORDER/CT
E7	0	2	NEURODEGENERATIVE DISORDERS/CT
E8	0	2.	NEURODERMATITIS/CT
E9	1		NEURODIA CYCLOPION FLORIDANA/CT
E10	0	2	NEUROECTODERM/CT
E11	0	2	NEUROECTODERM NEOPLASM/CT
E12	0	2	NEUROECTODERM TUMORS/CT

=> e e3+all E1 0 --> Neurodegenerative/CT ****** END ******

```
E# FREQUENCY AT
                                   TERM
                         ___
                         1 ALZHEIMER-TYPE/CT
2 ALZHEIMER-TYPE SENILE DEMENTIA/CT
           0
E1
                0
E2
               0 2 ALZHEIMER-TYPE
0 --> ALZHEIMER?/CT
0 1 AM/CT
0 2 AM 0.5/CT
0 2 AM 1002/CT
0 2 AM 100L/CT
0 2 AM 111/CT
0 2 AM 2000UP/CT
0 2 AM 6/CT
0 2 AM BLACK/CT
0 2 AM MICROBIAL 0
Е3
E4
E5
Ε6
E7
Ε8
E9
E10
E11
                                   AM MICROBIAL GENE AND GENETIC ELEMENT/CT
E12
=> e alzheimers/ct
E# FREQUENCY AT
                                   TERM
                          ___
                          1 ALZHEIMER-TYPE/CT
2 ALZHEIMER-TYPE SENILE DEMENTIA/CT
                0
E1
               0 1 ALZHEIMER-TYPE/CT
0 2 ALZHEIMER-TYPE SENILE DEMENTIA/CT
0 --> ALZHEIMERS/CT
0 1 AM/CT
0 2 AM 0.5/CT
0 2 AM 1002/CT
0 2 AM 100L/CT
0 2 AM 111/CT
0 2 AM 2000UP/CT
0 2 AM 6/CT
0 2 AM BLACK/CT
0 2 AM MICROBIAL GENE AND GENETIC ELEMENT/CT
E2
E3
E4
E5
Ε6
E7
E8
E9
E10
E11
E12
=> e cognitive impairment/ct
               O 2 COGNITIVE BEHAVIOR/CT
O 3 COGNITIVE DISORDERS/CT
O 3 --> COGNITIVE IMPAIRMENT/CT
O 3 COGNITIVE MENTAL DISORDER/CT
O 1 COGON/CT
O 2 COGON GRASS/CT
COGWHEELS/CT
E# FREQUENCY AT TERM
       -----
___
          0
F.1
E2
E3
E4
E5
Ε6
                COGWHEELS/CT
COH/CT
COHAERENS/CT
COHEN/CT
COHEN/CT
COHEN/CT
COHEN/CT
COHEN/CT
COHEN/CT
E7
E8
E9
E10
E11
E12
                   9
                                   COHEN, ERNST/CT
=> e e3+all
E.1
                        --> Cognitive impairment/CT
E2
                          USE Cognition (L) disorder/CT
                           USE Mental disorder (L) cognitive/CT
****** END ******
=> e neural trauma/ct
E# FREQUENCY AT
                                   TERM
                          ___
                                 NEURAL TISSUE-SPECIFIC F-ACTIN-BINDING PROTEINS/CT
                  0
E.1
                         3
                 0
0
E2
                                   NEURAL TRANSPLANT/CT
                           --> NEURAL TRAUMA/CT
Е3
Ε4
                          2 NEURAL TUBE/CT
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=> e alzheimer?/ct

```
0 2 NEURAL TUBE DEFECT/CT
0 2 NEURAL TUBE DEFECTS/CT
0 2 NEURAL TUBE DISEASE/CT
0 2 NEURAL TUBE NERVOUS SYSTEM/CT
0 2 NEURAL TUMORS/CT
0 1 NEURAL-/CT
E.5
E6
E7
E8
E9
E10
             0
                   2
                          NEURAL- AND THYMUS-DERIVED ACTIVATOR FOR ERBB KINASES/
E11
           55
                  2
E12
                          NEURALGIA/CT
=> fil medl, hcapl, embase, biosis; s l11 and (mental disorder(l) cognitive or
cognition(l)disorder)
COST IN U.S. DOLLARS
                                                    SINCE FILE
                                                                     TOTAL
                                                          ENTRY
                                                                   SESSION
FULL ESTIMATED COST
                                                           7.08
                                                                    735.96
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
                                                    SINCE FILE
                                                                    TOTAL
                                                         ENTRY SESSION
CA SUBSCRIBER PRICE
                                                           0.00
                                                                    -39.20
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FILE 'BIOSIS' ENTERED AT 10:38:28 ON 28 SEP 2004
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             O FILE MEDLINE
L17
L18
             O FILE HCAPLUS
L19
             O FILE EMBASE
L20
             O FILE BIOSIS
TOTAL FOR ALL FILES
L21
            0 L11 AND (MENTAL DISORDER(L) COGNITIVE OR COGNITION(L) DISORDER)
=> s rapin, j?/au;s witzmann, h?/au;s grumel, j?/au;s gonella, j?/au
          114 FILE MEDLINE
L23
           166 FILE HCAPLUS
L24
           131 FILE EMBASE
L25
           149 FILE BIOSIS
TOTAL FOR ALL FILES
L26
          560 RAPIN, J?/AU
L27
             3 FILE MEDLINE
L28
           134 FILE HCAPLUS
L29
             3 FILE EMBASE
L30
             3 FILE BIOSIS
TOTAL FOR ALL FILES
L31 143 WITZMANN, H?/AU
L32
             O FILE MEDLINE
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4 FILE HCAPLUS
              2 FILE EMBASE
L34
L35
              1 FILE BIOSIS
TOTAL FOR ALL FILES
L36
            7 GRUMEL, J?/AU
L37
             45 FILE MEDLINE
             38 FILE HCAPLUS
L38
L39
             29 FILE EMBASE
             47 FILE BIOSIS
L40
TOTAL FOR ALL FILES
            159 GONELLA, J?/AU
=> s 126 and 131 and 136 and 141
              O FILE MEDLINE
L43
              4 FILE HCAPLUS
L44
              0 FILE EMBASE
L45
              O FILE BIOSIS
TOTAL FOR ALL FILES
L46
              4 L26 AND L31 AND L36 AND L41
=> d 1-4 cbib abs; fill req
L46 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2004 ACS on STN
2002:609523 Document No. 137:155181 Synthesis of tripeptides and tripeptide
     derivatives for the treatment of neurodegenerative diseases. Rapin,
     Jean; Witzmann, Hans Klaus; Grumel, Jean-Marie;
     Gonella, Jacques (Tell-Pharm Ag, Switz.). Ger. Offen. DE 10105041
     Al 20020814, 12 pp. (German). CODEN: GWXXBX. APPLICATION: DE
     2001-10105041 20010205.
     The invention concerns the use of tripeptide derivs. [e.g.,
AB
     H-Gly-Phe-Pro-NH2 (I)] for the treatment of neurodegenerative disease,
     such as Alzheimer's disease. Thus, Boc-Phe-OH [Boc = (CH3)30C(0)] was coupled with TFA.H-Pro-NH2 to give a dipeptide, which was N-deprotected
     and converted to its TFA salt for coupling with Boc-Gly-OH; the resulting
     protected tripeptide was N-deprotected and converted to its HCl salt. The
     blood-brain partition coeffs. of I and seventeen similar tripeptides were
     given. The plasma half-life of 14C-labeled I.HCl was determined in rats (no data). Using a rat model of Alzheimer's disease, results of treatment
     with I showed retention of learned behavior in a five-day test of
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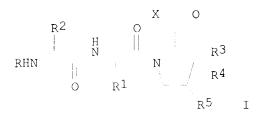
L46 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2004 ACS on STN 2002:609522 Document No. 137:163818 Tripeptide derivatives for the treatment of post-lesional diseases of the nervous system. Rapin, Jean; Witzmann, Hans Klaus; Grumel, Jean-Marie; Gonella, Jacques (Tell-Pharm AG, Switz.). Ger. Offen. DE 10105040 A1 20020814, 4 pp. (German). CODEN: GWXXBX. APPLICATION: DE 2001-10105040 20010205.

revealed increase dendrite development in the hippocampus.

pole-climbing at a signal to avoid shock. Examination of subject brains

GΙ

L33

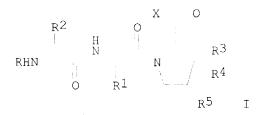


The invention discloses the use of cinnamoyl tripeptide derivs. for the treatment of post-lesional neuronal diseases. The cinnamoyl tripeptide derivs. are I [X = OH, C1-5 alkoxy, NH2, NH(C1-5 alkyl), N(C1-5 alkyl)2; R = (preferably) cinnamoyl; R1 = group derived from Phe, Tyr, Trp, Pro, Ala, Val, Leu or Ile; R2 = group derived from Gly, Ala, Ile, Val, Ser, Thr, His, Arg, Lys, Pro, Glu, Gln, pGlu, Asp and Asn; R3, R4 = H, OH, C1-5 alkyl, C1-5 alkoxy, provided that R3 and R4 are not both OH or C1-5 alkoxy; R5 = H, OH, C1-5 alkyl, C1-5 alkoxy], or a pharmaceutical acceptable salt thereof.

L46 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2004 ACS on STN
2002:609521 Document No. 137:140781 Synthesis of tripeptides and tripeptide derivatives for the treatment of post lesional diseases of the nervous system.. Rapin, Jean; Witzmann, Hans Klaus;
Grumel, Jean-Marie; Gonella, Jacques (Tell-Pharm Ag, Switz.). Ger. Offen. DE 10105038 A1 20020814, 10 pp. (German). CODEN: GWXXBX. APPLICATION: DE 2001-10105038 20010205.

AB The invention concerns the use of tripeptide derivs. [e.g., H-Gly-Phe-Pro-NH2 (I)] for the treatment of post lesional diseases of the nervous system. Thus, Boc-Phe-OH [Boc = (CH3)30C(0)] was coupled with TFA.H-Pro-NH2 to give a dipeptide, which was N-deprotected and converted to its TFA salt for coupling with Boc-Gly-OH; the resulting protected tripeptide was N-deprotected and converted to its HCl salt. The blood-brain partition coeffs. of I and seventeen similar tripeptides were given. The plasma half-life of 14C-labeled I.HCl was determined in rats (no data). Using an in-vivo dendritic sprouting assay in rats, I was tested for effect on hippocampus septum, and showed growth of up to $2\mu\text{m}$, compared to controls.

L46 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2004 ACS on STN
2002:591566 Document No. 137:135103 Tripeptide derivatives for treatment of neurodegenerative diseases. Rapin, Jean; Witzmann, Hans
Klaus; Grumel, Jean-Marie; Gonella, Jacques
(Tell-Pharm A.-G., Switz.). Ger. Offen. DE 10105039 A1 20020808, 10 pp.
(German). CODEN: GWXXBX. APPLICATION: DE 2001-10105039 20010205.



The invention discloses the use of tripeptide derivs. for treatment of neurodegenerative diseases. The tripeptide derivs. are I [X = OH, C1-5 alkoxy, NH2, NH(C1-5 alkyl), N(C1-5 alkyl)2; R = (preferably) cinnamoyl; R1 = group derived from Phe, Tyr, Trp, Pro, Ala, Val, Leu or Ile; R2 = group derived from Gly, Ala, Ile, Val, Ser, Thr, His, Arg, Lys, Pro, Glu, Gln, pGlu, Asp or Asn; R3, R4 = H, OH, C1-5 alkyl, C1-5 alkoxy, provided that R3 and R4 are not both OH or C1-5 alkoxy; R5 = H, OH, C1-5 alkyl, C1-5 alkoxy], or a pharmaceutically compatible salt. Cinnamoyl-Gly-L-Phe-L-Pro-NH2 was tested in an Alzheimer's disease model.

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	15.01	750.97
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
CA SUBSCRIBER PRICE	ENTRY -2.80	SESSION -42.00

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=> e	"cinnamoyl-gl	ycyl-l-phenylalanyl-l-prolinamide"/cn
El	1	CINNAMOYL-COA: (R)-PHENYLLACTATE COA-TRANSFERASE/CN
E2	1	CINNAMOYL-COENZYME A REDUCTASE/CN
ЕЗ	0>	CINNAMOYL-GLYCYL-L-PHENYLALANYL-L-PROLINAMIDE/CN
E4	1	CINNAMOYLACETALDEHYDE/CN
E5	1	CINNAMOYLACETONE/CN
E6	1	CINNAMOYLACETONITRILE/CN
E7	1	CINNAMOYLAUBUCIN/CN
E8	1	CINNAMOYLAUCUBIN/CN
E9	1	CINNAMOYLCHOLINE/CN
E10	1	CINNAMOYLCHOLINE CHLORIDE/CN
E11	1	CINNAMOYLCOCAINE/CN
E12	1	CINNAMOYLDIAMINOPIPERAZINE/CN
=>		
=> e		
E13	1	CINNAMOYLECGONINE/CN
E14	1	CINNAMOYLECGONINE METHYL ESTER/CN

```
CINNAMOYLETHYL METHACRYLATE-DIMETHYLOCTYLAMMONIUM STYRENESUL
E15
             1
                    FONATE-METHACRYLIC ACID-TRIS(TRIMETHYLSILOXY)(3-(METHACRYLOY
                    LOXY) PROPYL) SILANE COPOLYMER/CN
                    CINNAMOYLETHYL METHACRYLATE-STYRENE BLOCK COPOLYMER/CN
E16
             1
                    CINNAMOYLFERROCENE/CN
E17
             1
E18
             1
                   CINNAMOYLGLUCOSE-ALCOHOL CINNAMOYLTRANSFERASE/CN
                   CINNAMOYLGLYCINE/CN
E19
             1
                   CINNAMOYLGLYCINE P-NITROPHENYL ESTER/CN
E20
             1
                   CINNAMOYLGRANDIFLORIC ACID/CN
E21
             1
E22
             1
                   CINNAMOYLHYDRAZINE/CN
E23
             1
                   CINNAMOYLHYDROXAMIC ACID/CN
E24
             1
                   CINNAMOYLIUM/CN
=> s cinnamoyl(1)qlycyl(1)phenylalanyl(1)prolinamide
          3686 CINNAMOYL
             1 CINNAMOYLS
          3686 CINNAMOYL
                  (CINNAMOYL OR CINNAMOYLS)
        492293 GLYCYL
             2 GLYCYLS
        492293 GLYCYL
                  (GLYCYL OR GLYCYLS)
        367484 PHENYLALANYL
         21527 PROLINAMIDE
             O CINNAMOYL (L) GLYCYL (L) PHENYLALANYL (L) PROLINAMIDE
L47
=> e "cinnamoyl-isoleucyl-phenylalanyl-l-proline ethylamide"/cn
                   CINNAMOYL-COA: (R) - PHENYLLACTATE COA-TRANSFERASE/CN
                   CINNAMOYL-COENZYME A REDUCTASE/CN
Ε2
             1
             0 --> CINNAMOYL-ISOLEUCYL-PHENYLALANYL-L-PROLINE ETHYLAMIDE/CN
E3
                   CINNAMOYLACETALDEHYDE/CN
F.4
             7
E5
             1
                   CINNAMOYLACETONE/CN
E6
             1
                   CINNAMOYLACETONITRILE/CN
E7
             1
                   CINNAMOYLAUBUCIN/CN
E8
             1
                   CINNAMOYLAUCUBIN/CN
E9
             1
                   CINNAMOYLCHOLINE/CN
E10
             1
                   CINNAMOYLCHOLINE CHLORIDE/CN
             1
                   CINNAMOYLCOCAINE/CN
E11
             1
                   CINNAMOYLDIAMINOPIPERAZINE/CN
E12
=> s cinnamoyl(1)isoleucyl(1)phenylalanyl(1)proline(1)ethylamide
          3686 CINNAMOYL
             1 CINNAMOYLS
          3686 CINNAMOYL
                  (CINNAMOYL OR CINNAMOYLS)
        325112 ISOLEUCYL
        367484 PHENYLALANYL
         85965 PROLINE
             2 PROLINES
         85965 PROLINE
                  (PROLINE OR PROLINES)
           621 ETHYLAMIDE
L48
             O CINNAMOYL(L) ISOLEUCYL(L) PHENYLALANYL(L) PROLINE(L) ETHYLAMIDE
=> e "cinnamoyl-isoleucyl-isoleucyl-prolineamide"/cn
E.1
                   CINNAMOYL-COA: (R) - PHENYLLACTATE COA-TRANSFERASE/CN
                   CINNAMOYL-COENZYME A REDUCTASE/CN
E2
E3
             0 --> CINNAMOYL-ISOLEUCYL-ISOLEUCYL-PROLINEAMIDE/CN
E4
                   CINNAMOYLACETALDEHYDE/CN
             1
E5
             1
                   CINNAMOYLACETONE/CN
Ε6
             1
                   CINNAMOYLACETONITRILE/CN
E7
             1
                   CINNAMOYLAUBUCIN/CN
```

```
E8
             1
                   CINNAMOYLAUCUBIN/CN
E9
             1
                   CINNAMOYLCHOLINE/CN
E10
             1
                   CINNAMOYLCHOLINE CHLORIDE/CN
E11
             1
                   CINNAMOYLCOCAINE/CN
E12
             1
                   CINNAMOYLDIAMINOPIPERAZINE/CN
=> s cinnamoyl(l)isoleucyl(l)isoleucyl(l)prolineamide
          3686 CINNAMOYL
             1 CINNAMOYLS
          3686 CINNAMOYL
                  (CINNAMOYL OR CINNAMOYLS)
        325112 ISOLEUCYL
        325112 ISOLEUCYL
             2 PROLINEAMIDE
             O CINNAMOYL (L) ISOLEUCYL (L) ISOLEUCYL (L) PROLINEAMIDE
L49
=> fil medl, hcap, biosis, embase, wpids
COST IN U.S. DOLLARS
                                                  SINCE FILE
                                                                  TOTAL
                                                       ENTRY
                                                              SESSION
FULL ESTIMATED COST
                                                       59.69
                                                                810.66
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)
                                                  SINCE FILE
                                                                  TOTAL
                                                      ENTRY
                                                               SESSION
CA SUBSCRIBER PRICE
                                                       0.00
                                                                 -42.00
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=> s cinnamoyl(l)isoleucyl(l)isoleucyl(l)prolineamide
L50
            O FILE MEDLINE
L51
             0 FILE HCAPLUS
L52
            0 FILE BIOSIS
L53
             O FILE EMBASE
L54
             O FILE WPIDS
TOTAL FOR ALL FILES
            O CINNAMOYL(L) ISOLEUCYL(L) ISOLEUCYL(L) PROLINEAMIDE
=> s cinnamoyl(1)isoleucyl(1)phenylalanyl(1)proline(1)ethylamide
          O FILE MEDLINE
L57
             0 FILE HCAPLUS
L58
             0 FILE BIOSIS
L59
            O FILE EMBASE
L60
            O FILE WPIDS
TOTAL FOR ALL FILES
L61
            0 CINNAMOYL(L) ISOLEUCYL(L) PHENYLALANYL(L) PROLINE(L) ETHYLAMIDE
=> s cinnamoyl(1)glycyl(1)phenylalanyl(1)prolinamide
```

O FILE MEDLINE L62 L63 O FILE HCAPLUS 0 FILE BIOSIS L64 L65O FILE EMBASE 1 FILE WPIDS L66 TOTAL FOR ALL FILES 1 CINNAMOYL(L) GLYCYL(L) PHENYLALANYL(L) PROLINAMIDE => d L67 ANSWER 1 OF 1 WPIDS COPYRIGHT 2004 THE THOMSON CORP on STN 2002-667977 [72] ΑN WPIDS DNC C2002-187773 Use of proline-containing tripeptide for treating neurodegeneration, TΤ useful particularly in early stages of Alzheimer's disease. DC B02 B03 ΙN GONELLA, J; GRUMEL, J; RAPIN, J; WITZMANN, H K (TELL-N) TELL-PHARM AG; (NEUR-N) NEUROTELL AG PΑ CYC 101 PΙ DE 10105039 A1 20020808 (200272)* A61K038-06 10 WO 2002062830 A1 20020815 (200272) EN C07K005-08 RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ NL OA PT SD SE SL SZ TR TZ UG ZM ZW W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ UA UG US UZ VN YU ZA ZM ZWEP 1358204 A1 20031105 (200377) EN C07K005-08 R: AL AT BE CH CY DE DK ES FI FR GB GR IE IT LI LT LU LV MC MK NL PT RO SE SI TR AU 2002247685 A1 20020819 (200427) C07K005-08 DE 10105039 A1 DE 2001-10105039 20010205; WO 2002062830 A1 WO 2002-EP1181 ADT 20020205; EP 1358204 A1 EP 2002-716727 20020205, WO 2002-EP1181 20020205; AU 2002247685 A1 AU 2002-247685 20020205 FDT EP 1358204 Al Based on WO 2002062830; AU 2002247685 Al Based on WO 2002062830 PRAI DE 2001-10105039 20010205 ICM A61K038-06; C07K005-08 ICS A61P025-28 => fil reg COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 11.98 822.64 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE 0.00 -42.00 FILE 'REGISTRY' ENTERED AT 10:43:58 ON 28 SEP 2004 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2004 American Chemical Society (ACS)

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